

EDITORIAL BOARD

EDITOR IN CHIEF

Professor G I Muguti

ASSOCIATE EDITORS

Professor IT Gangaidzo

Dr S P Munjanja

EDITORIAL BOARD MEMBERS

<i>Professor MM Chidzonga</i>	<i>(Zimbabwe)</i>
<i>Professor P Jacobs</i>	<i>(South Africa)</i>
<i>Dr R A Kambarami</i>	<i>(Zimbabwe)</i>
<i>Professor A S Latif</i>	<i>(Zimbabwe)</i>
<i>Professor P R Mason</i>	<i>(Zimbabwe)</i>
<i>Professor CT Musabayane</i>	<i>(Zimbabwe)</i>
<i>Professor KJ Nathoo</i>	<i>(Zimbabwe)</i>
<i>Mr L Nystrom</i>	<i>(Sweden)</i>
<i>Dr S Siziya</i>	<i>(Zambia)</i>

PAST EDITORS

Professor M Gelfand (1953-1985)

Professor H M Chinyanga (1985-1990)

Professor J A Matenga (1991-1999)

ADMINISTRATIVE AND OFFICE STAFF

Director of Publications: Mr Munani S Mtetwa

Administrative Manager: Mr Christopher Mashavira

Technical Editor: Mrs Ling M Cooper

Statistical Advisor: Mr S Rusakaniko

Secretary: Mrs Patricia Bhunhu

All manuscripts will be prepared with the International Committee of Medical Journal Editors - Uniform requirements for manuscripts submitted to Biomedical Journals, 1993.

Manuscripts submitted for publication are accepted on the understanding that they are contributed exclusively to *The Central African Journal of Medicine*. A statement to that effect should be included in the letter accompanying the manuscript.

Communications concerning editorial matter, advertising, subscriptions, change of address, etc. Should be addressed to the Administrative Manager, P. O. Box A195 Avondale, Harare, Zimbabwe.

The subscription rate for **surface transmission** including postage for year 2002 is Z\$3 100.00 locally; Africa US\$210.00 for individuals and US\$265.00 for institutions; and US\$260.00 for individuals and US\$280.00 for institutions for the rest of the world per annum. The subscription rate for **airmail transmission** for year 2002 in Africa is US\$300.00 for individuals US\$320.00 for institutions and US\$70.00 for postage; and US\$330.00 for individuals US\$350.00 for institutions and US\$70.00 for postage for the rest of the world per annum.

Owned and published by the Central African Journal of Medicine in conjunction with the Faculty of Medicine



University of Zimbabwe

The influence of socio-economic factors on *Helicobacter pylori* infection rates of students in rural Zambia

*NJD MCLAUGHLIN, **DI MCLAUGHLIN, *H LEFCORT

Abstract

Objectives: Although prevalence of disease in sub-Saharan Africa is often quite high and attracts much research, relatively little is known about less critical maladies. We examined *Helicobacter pylori* infected students in rural Zambia. We attempted to determine if any socio-economic or co-occurring diseases were correlated to *H. pylori* infection. Understanding the context in which *H. pylori* infections occur may increase our understanding of this organism.

Design: We conducted a screening survey with diagnostic tests of primary and secondary school students to determine rates of *H. pylori* infection. We then correlated these rates to socio-economic factors such as income and tobacco use. We also explored the correlation of *H. pylori* to HIV and malaria.

Setting: Zimba, Zambia.

Subjects: Eighty seven primary and secondary school students.

Main Outcome Measure: Correlation of *H. pylori* to socio-economic factors.

Results: *H. pylori* infection was common (60.9%) and was consistent with rates found in other African countries. We found no significant correlation between *H. Pylori* and disease and socio-economic variables.

Conclusion: In the studied population *H. pylori* infection does not appear to be correlated with the measured socio-economic or disease variables.

Cent Afr J Med 2003;49(3/4):38-41

Introduction

People in rural sub-Saharan Africa face the interrelated problems of high disease rates, high poverty, and poor access to health care. Diseases such as malaria and Acquired Immunodeficiency Syndrome (AIDS) are widespread, and in many areas are increasing.¹ Furthermore, lack of access to medical resources precludes care for all except in the final stages of disease.

Compounding these problems is a second tier of less severe maladies such as chronic anaemia and colonization with the gastric bacterium *Helicobacter pylori* which is associated with gastric and duodenal ulcers. Although less pressing, these other maladies may interact and affect the morbidity and mortality of diseases such as AIDS and malaria.

Other factors contributing to poor health are socio-economic variables such as income, education, the number of sexual partners, and tobacco use. For example, some

studies indicate a positive correlation between *H. pylori* and tobacco smoking^{2,3} although other studies either do not support this⁴ or have even found a negative correlation.^{5,6} Similarly, income and education have been shown to be negatively correlated with *H. pylori* rates in Scotland³ and Ethiopia,⁷ although other studies in South Africa⁸ and Senegal⁹ have not found this. Furthermore, due to cultural practices, *H. pylori* is correlated to religious affiliation and urban habitation when other socio-economic factors are controlled.⁷

H. pylori is widespread in Africa. Infection rates for adults are 85% in Nigeria,¹⁰ 82.8% in Senegal,⁹ and 77.5% in Zaire.¹¹ In a review of published studies Kidd et al¹² found that throughout Africa about 61% of all adults test positive for *H. pylori* but most are asymptomatic.

H. pylori also appears to be contracted at a young age. In Nigeria 82% of those infected acquire the bacterium by age 10;¹⁰ in South Africa 67.3%¹³ and 80%⁸ by age 10; and in Zaire 66% by age nine.¹¹

*Biology Department
Gonzaga University
Spokane, WA
99258
USA

**The Bozeman Clinic
931 Highland Blvd Suite 3360
Bozeman, MT
59715
USA

Correspondence to:
Dr Hugh Lefcort
e-mail: lefcort@gonzaga.edu
TEL: 509-323-6706
FAX: 509-323-5804

Although prevalence of disease in sub-Saharan Africa is often quite high, relatively little is known about associated health and social demographics. We examined students and determined rates of *H. pylori* and then we explored if correlations exist between *H. pylori* and rates of infection with diseases and to socio-economic factors. Understanding the context in which *H. pylori* infections occur may increase our understanding of this organism.

Materials and Methods

Background.

The study was conducted between 19 January 2000 and 19 February 2000 in Zimba, Zambia. Zimba has a population of 4 000 people and is located in rural southwestern Zambia about 70 km north of Livingstone. Most of the people are employed in subsistence agriculture (maize, ground nuts, and cabbage) but a small percentage work as teachers, nurses, and low level government officials. Water is drawn from bore holes and a nearby lake. Health care is limited and is mainly provided by state clinics and a missionary hospital. Maladies of protein deficiency such as Kwashiorkor are as widespread as is HIV. Tuberculosis rates have also experienced a recent increase.

Selection of Subjects.

We surveyed 87 students (40 females and 47 males, aged 14 to 23) at three local schools. One school was a primary day school made up of poor students, one was a technical school of slightly higher income students, and one was a secondary boarding school of more affluent pupils.

An announcement of free screening was made once to all classes that were meeting on one given day. Students readily volunteered and every student in each of the classes we chose went to the clinic. The students were not compensated. Donors were anonymous (given a number and not asked their names) and individual test results were provided if requested.

Parental permission was obtained and ethical approval was given by the Gonzaga University Animal Care and Ethics Committee (12/10/99), the Zimba Mission Hospital Director, and each school Headmaster.

We conducted individual oral interviews and asked the students about age, gender, town of origin, tobacco use number of lifetime sexual partners, past treatment for malaria and present abdominal pain.

We also determined degree of poverty/wealth by asking the students about the occupation of their parents.¹⁴ *Poor* was defined as both parents engaged in subsistence agriculture. *Moderate* as one parent or member in the household working as a nurse, teacher, government official, and/or attending the technical school and *affluent* as able to attend the secondary boarding school. These categories and rankings were confirmed by the class teachers in all cases except one and that student was excluded from the analysis. An attempt was made during data analysis to control for family and extended family size but the different income categories did not differ for this variable.

Parameters such as "past treatment of malaria", and "present abdominal pain" are subjective since fever and/or headache is sometimes referred to as malaria in this geographic region. Additionally "reported abdominal pain" is similarly broad since it may include gynaecological symptoms in adolescent females. However, since we were dealing with children our choice of terms was intended to be subjective. The questions were asked only to establish a general state of health. The actual diagnosis of *H. Pylori*, HIV, or malaria was done using the fluid samples we collected.

Laboratory Tests.

At the Zimba Mission Hospital we drew two millilitres of blood from each student which we tested within 12 hours for HIV, malaria, *H. pylori*, and haematocrit. Blood was drawn by syringe from the median cubital vein and then injected into both a citrated capillary tube and a noncitrated capillary tube. Malaria infection was determined using Field A and B stains and rated by severity as 1+, 2+, and 3+, with 1+ being considered mild and 2+ and 3+ being severe. *H. pylori* was determined using Quickview tests (Quidel Corporation) which is a lateral-flow immunoassay that detects IgG antibodies specific to *H. pylori*. Haematocrit was determined by spinning the capillary tubes and then using a haematocrit reader (Clay Adams Co.). HIV was tested using a Western Blot test (Astra Pharmaceuticals) of blood plasma

Statistical Tests.

We used parametric Pearson Product Moment Correlations (PPMC) for continuous variables or variables with more than three categories and nonparametric Kruskal Wallis One Way ANOVA on Ranks for data with only two categories. Alpha was set to 0.05.

Results

H. pylori was present in 60.9% of the students and it was correlated with lower haematocrit levels (PPMC = -0.250, $n = 87$, $p = 0.019$, Figure I). Haematocrit was also lower in females (Kruskal Wallis One Way ANOVA, $H = 4.45$, d.f. = 1, $p = 0.035$) although it was consistent with the normal drop due to menses seen in females when compared to males. *H. pylori* was lower in students who smoked cigarettes but this did not reach statistical significance (Kruskal Wallis One Way ANOVA, $H = 3.42$, d.f. = 1, $p = 0.064$, Figure II).

Students who smoked reported greater abdominal pain (PPMC = 0.281, $n = 87$, $p = 0.011$) and more sexual partners (PPMC = 0.396, $n = 87$, $p = 0.0003$). Smoking rates were also higher in older age groups with 19 to 23 year olds smoking more than 14 to 18 year olds (Kruskal Wallis One Way ANOVA, $H = 3.93$, d.f. = 1, $p = 0.047$).

The HIV positive rate among students was 9.2%. Infection with HIV was positively correlated with age (PPMC = 0.303, $n = 87$, $p = 0.004$, Figure III) while *H. pylori* infection was slightly, but not significantly, correlated with age (PPMC = 0.166, $n = 87$, $p = 0.125$, Figure IV). Only one

Figure I: Percentage haematocrit levels of those with and without *H. pylori* infections.

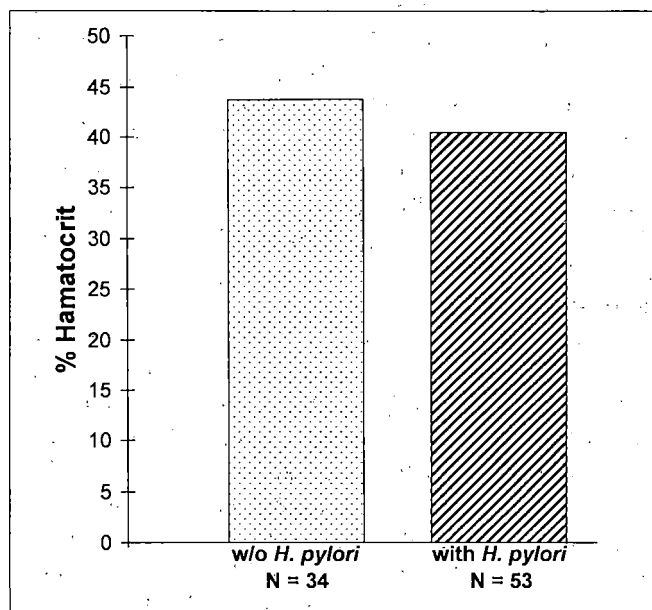
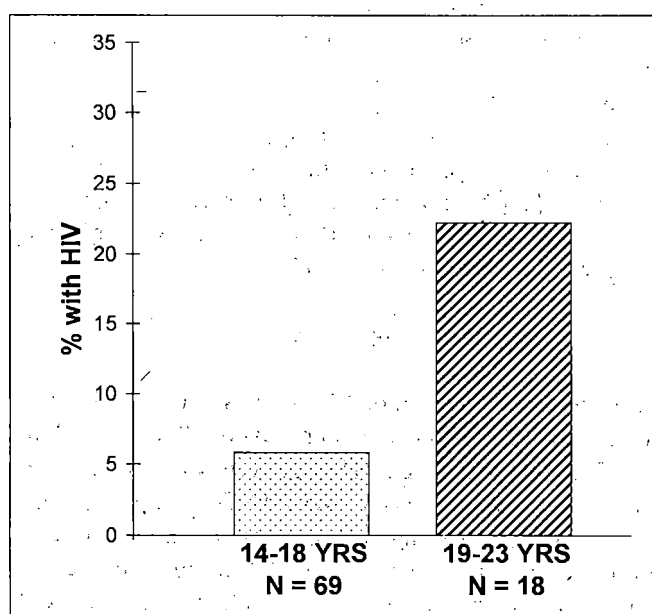


Figure III: HIV infection rates of younger and older students.



student had an active case of malaria. The lack of other active malaria infections was due to the schools' policies of sending ill students home. However, 77.8% of the students reported past treatment for malaria.

All other possible combinations of variables were not significantly correlated.

Discussion

H. pylori infection was common (60.9%) although it was consistent with rates found in other African countries. *H. pylori* was not statistically correlated with any socio-economic or disease variable that we measured.

Figure II: *H. pylori* rates of smokers and nonsmokers of tobacco cigarettes.

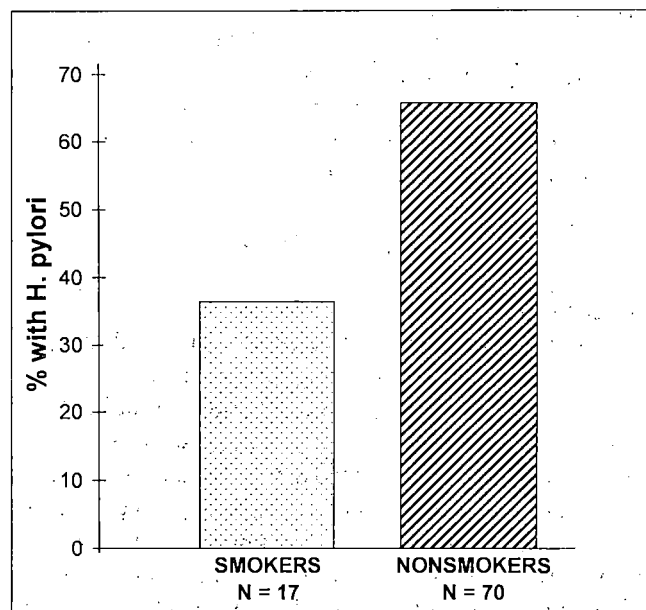
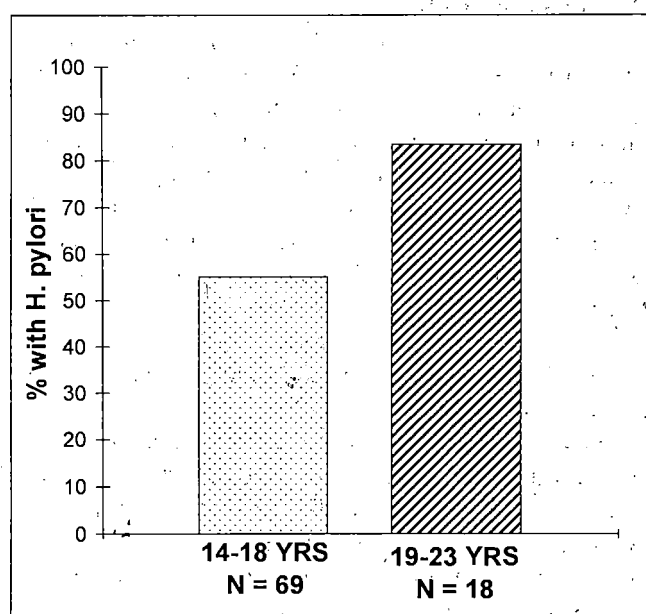


Figure IV: *H. pylori* infection rates of younger and older students.



We found that *H. pylori* was not correlated to self-reported gastro-intestinal upset. *H. pylori* is often associated with gastro-intestinal upset and ulceration.^{12,15} In Zairians with gastroduodenal disorders, *H. pylori* has been found in 87.5% of patients,¹¹ while another study found rates of 91.7%.¹⁶ However, it is not correlated with self-reported dyspepsia in the United Kingdom¹⁷ or Australia.¹⁸ Studies in more affluent countries like Taiwan² and Australia¹⁸ seem to suggest that the debilitating effects of *H. pylori* are higher in men, but studies in Africa have not found this.^{8,9,19} We found equal rates of *H. pylori* in both males and females and both reported equal rates of gastro-intestinal upset (39.6%).

H. pylori was correlated with lower haematocrit levels but it is unclear from our correlational data whether *H. pylori* is the cause of lower haematocrit or whether people who are partially anaemic are more susceptible to *H. pylori*.

There has been much debate on the association between *H. pylori* and smoking tobacco. Some studies have found a positive correlation^{2,3} while others find no correlation,⁴ or a negative correlation.^{5,6} We found a nearly significant ($p = 0.064$) negative correlation between smoking and *H. pylori*. Ogihara *et al*⁶ argue that since smoking increases gastric acidity it may defend against *H. pylori*. Yet increased gastric acidity could also cause increased peptic ulcers. Indeed, smokers in our study reported significantly greater rates of abdominal pain than non smokers.

In conclusion, in the studied population *H. pylori* infection does not appear to be correlated with the measured socioeconomic or disease variables.

Acknowledgements

We wish to thank World Medical Missions for arranging our visit and D Mwafe for laboratory services. We would also like to thank D Boose, D Lefcort, and S Palpant for reading earlier drafts of this manuscript. Funding was provided in part by Astra Pharmaceuticals.

References

1. Cohen J. Ground zero: AIDS research in Africa. *Science* 2000;288:2150-3.
2. Chen TS, Chang FY, Lee SD. Smoking and male gender rather than CagA protein are associated with increased risk for duodenal ulcer in *Helicobacter pylori* infected patients in Taiwan. *Dig Dis Sc* 1999;44:2076-80.
3. Woodward M, Morrison C, McColl K. An investigation into factors associated with *Helicobacter pylori* infection. *J Clin Epidemiol* 2000;53:175-81.
4. Holcombe C, Kaluba J, Lucas SB. Non-ulcer dyspepsia in Nigeria: a case-control study. *Trans R Soc Trop Med Hyg* 1991;85:553-5.
5. Brenner H, Rothenbacher D, Bode G, Adler G. Relation of smoking and alcohol and coffee consumption to active *Helicobacter pylori* infection: cross sectional study. *B M J* 1997;315(7121):1489-92.
6. Ogihara A, Kikuchi S, Hasegawa A, Kurosawa M, Miki K, Kaneko E, Mizukoshi H. Relationship between *Helicobacter pylori* infection and smoking and drinking habits. *J Gastroenterol Hepatol* 2000;15:271-6.
7. Lindkvist P, Enquselassie R, Asrat D, Muhe L, Nilsson I, Giesecke J. Risk factors for infection with *Helicobacter pylori* — a study of children in rural Ethiopia. *Scand J Infect Dis* 1998;30:371-6.
8. Sathar MA, Gouws E, Simjee AE, Mayat AM. Seroepidemiological study of *Helicobacter pylori* infection in South African children. *Trans R Soc Trop Med Hyg* 1997;91:393-5.
9. Mbengue M, Diouf ML, Dangou JM, Ka MM, Ba-Seck A, Ndiaye MF, *et al*. Frequency of *Helicobacter pylori* infection in symptomatic patients in Senegal. *Med Trop (Mars)* 1997;57:256-8.
10. Holcombe C, Omotara BA, Eldridge J, Jones DM. *H. pylori*, the most common bacterial infection in Africa: a random serological study. *Am J Gastroenterol* 1992;87:28-30.
11. Glupczynski Y, Bourdeaux L, Verhas M, DePrez C, DeVos D, Devreker D. Use of a urea breath test versus invasive methods to determine the prevalence of *Helicobacter pylori* in Zaire. *Eur J Clin Microbiol Infect Dis* 1992;11:322-7.
12. Kidd M, Louw JA, Marks IN. *Helicobacter pylori* in Africa: observations on an 'enigma within an enigma'. *J Gastroenterol Hepatol* 1999;14:851-8.
13. Pelsers HH, Househam KC, Joubert G, van der Linde G, Kraaij P, Meinardi, *et al*. Prevalence of *Helicobacter pylori* antibodies in children in Bloemfontein, South Africa. *J Paediatr Gastroenterological Nutrition* 1997;24:135-9.
14. Mazumdar, K. Classification of countries: a socio-economic approach. *Social Indicators Res* 1995;34:261-73.
15. Holcombe C, Lucas SB, Umar H, Abba A. *Helicobacter* (= *Campylobacter*) *pylori* in Africa. *Trans R Soc Trop Med Hyg* 1990;84:294-6.
16. Watters DA, Gilmour HM. *Helicobacter pylori* and non-ulcer dyspepsia in Zambia. *Trop Doctor* 1992;22:85.
17. Stone MA, Barnett DB, Mayberry JF. Lack of correlation between self-reported symptoms of dyspepsia and infection with *Helicobacter pylori*, in a general population sample. *Eur J Gastroenterol Hepatol* 1998;10:301-4.
18. Nandurkar S, Talley NJ, Xia H, Mitchell H, Hazel S, Jones M. Dyspepsia in the community is linked to smoking and aspirin use but not to *Helicobacter pylori* infection. *Arch Int Med* 1998;158:1427-33.
19. Eurogast Study Group. Epidemiology of, and risk factors for, *Helicobacter pylori* infection among 3 194 asymptomatic subjects in 17 populations. *Gut* 1993;34:1672-6.



This work is licensed under a
Creative Commons
Attribution – NonCommercial - NoDerivs 3.0 License.

To view a copy of the license please see:
<http://creativecommons.org/licenses/by-nc-nd/3.0/>

This is a download from the BLDS Digital Library on OpenDocs
<http://opendocs.ids.ac.uk/opendocs/>